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NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2	Apr 08	"Ask CAS" for self-help around the clock
NEWS	3	Apr 09	BEILSTEIN: Reload and Implementation of a New Subject Area
NEWS	4	Apr 09	ZDB will be removed from STN
NEWS	5	Apr 19	US Patent Applications available in IFICDB, IFIPAT, and IFIUDB
NEWS	6	Apr 22	Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS
NEWS	7	Apr 22	BIOSIS Gene Names now available in TOXCENTER
NEWS	8	Apr 22	Federal Research in Progress (FEDRIP) now available
NEWS	9	Jun 03	New e-mail delivery for search results now available
NEWS	10	Jun 10	MEDLINE Reload
NEWS	11	Jun 10	PCTFULL has been reloaded
NEWS	12	Jul 02	FOREGE no longer contains STANDARDS file segment
NEWS	13	Jul 22	USAN to be reloaded July 28, 2002; saved answer sets no longer valid
NEWS	14	Jul 29	Enhanced polymer searching in REGISTRY
NEWS	15	Jul 30	NETFIRST to be removed from STN
NEWS	16	Aug 08	CANCERLIT reload
NEWS	17	Aug 08	PHARMAMarketLetter(PHARMAML) - new on STN
NEWS	18	Aug 08	NTIS has been reloaded and enhanced
NEWS	19	Aug 19	Aquatic Toxicity Information Retrieval (AQUIRE) now available on STN
NEWS	20	Aug 19	IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS	21	Aug 19	The MEDLINE file segment of TOXCENTER has been reloaded
NEWS	22	Aug 26	Sequence searching in REGISTRY enhanced
NEWS	23	Sep 03	JAPIO has been reloaded and enhanced
NEWS	24	Sep 16	Experimental properties added to the REGISTRY file
NEWS	25	Sep 16	CA Section Thesaurus available in CAPLUS and CA
NEWS	26	Oct 01	CASREACT Enriched with Reactions from 1907 to 1985
NEWS	27	Oct 21	EVENTLINE has been reloaded
NEWS	28	Oct 24	BEILSTEIN adds new search fields
NEWS	29	Oct 24	Nutraceuticals International (NUTRACEUT) now available on STN
NEWS	30	Oct 25	MEDLINE SDI run of October 8, 2002
NEWS	31	Nov 18	DKILIT has been renamed APOLLIT
NEWS	32	Nov 25	More calculated properties added to REGISTRY
NEWS	33	Dec 02	TIBKAT will be removed from STN
NEWS	34	Dec 04	CSA files on STN
NEWS	35	Dec 17	PCTFULL now covers WP/PCT Applications from 1978 to date
NEWS	36	Dec 17	TOXCENTER enhanced with additional content
NEWS	37	Dec 17	Adis Clinical Trials Insight now available on STN
NEWS	38	Dec 30	ISMEC no longer available
NEWS	39	Jan 13	Indexing added to some pre-1967 records in CA/CAPLUS
NEWS	40	Jan 21	NUTRACEUT offering one free connect hour in February 2003
NEWS	41	Jan 21	PHARMAML offering one free connect hour in February 2003
NEWS	42	Jan 29	Simultaneous left and right truncation added to COMPENDEX, ENERGY, INSPEC
NEWS	43	Feb 13	CANCERLIT is no longer being updated
NEWS	44	Feb 24	METADDEX enhancements
NEWS	45	Feb 24	PCTGEN now available on STN

NEWS 46 Feb 24 TEMA now available on STN  
NEWS 47 Feb 26 NTIS now allows simultaneous left and right truncation  
NEWS 48 Feb 26 PCTFULL now contains images

NEWS EXPRESS January 6 CURRENT WINDOWS VERSION IS V6.01a,  
CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),  
AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002  
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NEWS INTER General Internet Information  
NEWS LOGIN Welcome Banner and News Items  
NEWS PHONE Direct Dial and Telecommunication Network Access to STN  
NEWS WWW CAS World Wide Web Site (general information)

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=> file .gary

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'MEDLINE' ENTERED AT 11:11:19 ON 04 MAR 2003

FILE 'CANCERLIT' ENTERED AT 11:11:19 ON 04 MAR 2003

FILE 'BIOSIS' ENTERED AT 11:11:19 ON 04 MAR 2003

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FILE 'SCISEARCH' ENTERED AT 11:11:19 ON 04 MAR 2003

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=> s (kozarov-E? or progulske-a? or progulske-fox-a?)/au

L1 226 (KOZAROV-E? OR PROGULSKE-A? OR PROGULSKE-FOX-A?)/AU

=> s l1 and gingivalis

L2 112 L1 AND GINGIVALIS

=> s l2 and angio? or neovas?

L3 58378 L2 AND ANGIO? OR NEOVAS?

=> s l2 and (angio? or neovas?)

L4 0 L2 AND (ANGIO? OR NEOVAS?)

=> s l2 and (?carcinoma or cancer? or neoplas? or tumor? or metast?)

2 FILES SEARCHED...

L5 0 L2 AND (?CARCINOMA OR CANCER? OR NEOPLAS? OR TUMOR? OR METAST?)

=> s l2 and (HagA or Hemag?)  
L6 68 L2 AND (HAGA OR HEMAG?)

=> dup rem l6  
PROCESSING COMPLETED FOR L6  
L7 38 DUP REM L6 (30 DUPLICATES REMOVED)

=> s l7 and anti?  
2 FILES SEARCHED...  
L8 13 L7 AND ANTI?

=> d ibib abs 1-38 l7

L7 ANSWER 1 OF 38 SCISEARCH COPYRIGHT 2003 ISI (R)  
ACCESSION NUMBER: 2001:396093 SCISEARCH  
THE GENUINE ARTICLE: 430HD  
TITLE: The effect of monoclonal antibody and route of immunization on the humoral immune response against *Porphyromonas gingivalis*  
AUTHOR: van Tilburg M L J A; Kozarov E V; Progulske-Fox A; Brady L J (Reprint)  
CORPORATE SOURCE: Univ Florida, Dept Oral Biol, JHMH Box 100424, Gainesville, FL 32610 USA (Reprint); Univ Florida, Dept Oral Biol, Gainesville, FL 32610 USA  
COUNTRY OF AUTHOR: USA  
SOURCE: ORAL MICROBIOLOGY AND IMMUNOLOGY, (JUN 2001) Vol. 16, No. 3, pp. 153-162.  
Publisher: MUNKSGAARD INT PUBL LTD, 35 NORRE SOGADE, PO BOX 2148, DK-1016 COPENHAGEN, DENMARK.  
ISSN: 0902-0055.  
DOCUMENT TYPE: Article; Journal  
LANGUAGE: English  
REFERENCE COUNT: 51

\*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS\*

AB Immunomodulation mediated by exogenous antibodies has been proposed as a vaccine strategy to improve immune protection against pathogenic microorganisms and suggested to contribute to protection following passive immunization. To test whether a monoclonal antibody directed against an adhesion epitope of the periodontal pathogen *Porphyromonas gingivalis* could influence the humoral immune response following mucosal immunization, BALB/c mice were immunized orally or intranasally with P. *gingivalis* alone or P. *gingivalis* coated with monoclonal antibody 61BG1.3. Differences in antigenic specificity of anti-P. *gingivalis* serum immunoglobulin G (IgG) were demonstrated between groups of mice that received monoclonal antibody-coated P. *gingivalis* versus those that received P. *gingivalis* alone by either route of immunization. Binding of monoclonal antibody 61BG1.3 to P. *gingivalis* prior to immunization did not influence the serum IgG subclass distribution. However, minor differences in subclass distribution were observed between the various routes of mucosal immunization. These results support the hypothesis that specific monoclonal antibody bound to a bacterial vaccine can alter the quality of the humoral immune response to that organism.

L7 ANSWER 2 OF 38 MEDLINE DUPLICATE 1  
ACCESSION NUMBER: 2000316075 MEDLINE  
DOCUMENT NUMBER: 20316075 PubMed ID: 10858264  
TITLE: Long-term immunological memory induced by recombinant oral *Salmonella* vaccine vectors.  
AUTHOR: Kohler J J; Pathangey L; Hasona A; Progulske-Fox A ; Brown T A  
CORPORATE SOURCE: Department of Oral Biology, University of Florida, Gainesville, Florida 32610, USA.  
CONTRACT NUMBER: DE-07200 (NIDCR)

DE-07496 (NIDCR)

DE-10963 (NIDCR)

SOURCE: INFECTION AND IMMUNITY, (2000 Jul) 68 (7) 4370-3.  
Journal code: 0246127. ISSN: 0019-9567.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 200007  
ENTRY DATE: Entered STN: 20000728  
Last Updated on STN: 20000728  
Entered Medline: 20000720

AB We have previously shown that *Salmonella enterica* serovar Typhimurium expressing the *hagB* **hemagglutinin** gene from *Porphyromonas gingivalis* can induce primary and recall immune responses in serum and secretions in mice; however, the longevity of memory induced by oral *Salmonella* carriers has not been adequately demonstrated. In this study, we examined the capacity of mice to mount a recall response 52 weeks after primary immunization. Recall responses were seen in serum immunoglobulin G (IgG) and IgA following boosting at week 52, and in most cases, they were equal to or greater than the primary responses. Significant mucosal IgA recall responses in saliva and vaginal wash were also detected following boosting at week 52. In addition, there was a considerable residual response in secretions at week 51, prior to boosting. These results indicate that oral *Salmonella* vectors can induce long-term memory to recombinant *HagB* and are particularly effective at inducing long-lasting mucosal responses as well as at inducing the capacity for mucosal recall responses.

L7 ANSWER 3 OF 38 SCISEARCH COPYRIGHT 2003 ISI (R)  
ACCESSION NUMBER: 2000:164400 SCISEARCH  
THE GENUINE ARTICLE: 277MH  
TITLE: Expression, purification, immunological and functional properties of recombinant **hemagglutinin A** from *Porphyromonas gingivalis*  
AUTHOR: **Kozarov E (Reprint)**; Nowacki C; Progulske-Pox A  
CORPORATE SOURCE: UNIV FLORIDA, DEPT ORAL BIOL, GAINESVILLE, FL 32610  
COUNTRY OF AUTHOR: USA  
SOURCE: JOURNAL OF DENTAL RESEARCH, (15 FEB 2000) Vol. 79, Sp. iss. SI, pp. 2005-2005.  
Publisher: AMER ASSOC DENTAL RESEARCH, 1619 DUKE ST, ALEXANDRIA, VA 22314.  
ISSN: 0022-0345.  
DOCUMENT TYPE: Conference; Journal  
FILE SEGMENT: LIFE; CLIN  
LANGUAGE: English  
REFERENCE COUNT: 0

L7 ANSWER 4 OF 38 MEDLINE DUPLICATE 2  
ACCESSION NUMBER: 2000107078 MEDLINE  
DOCUMENT NUMBER: 20107078 PubMed ID: 10639440  
TITLE: Expression and immunogenicity of **hemagglutinin A** from *Porphyromonas gingivalis* in an avirulent *Salmonella enterica* serovar typhimurium vaccine strain.  
AUTHOR: **Kozarov E**; Miyashita N; Burks J; Cervený K; Brown T A; McArthur W P; **Progulske-Fox A**  
CORPORATE SOURCE: Department of Oral Biology and the Periodontal Disease Research Center, University of Florida, Gainesville, Florida 32610, USA.. kozarov@dental.ufl.edu  
CONTRACT NUMBER: DE07496 (NIDCR)  
SOURCE: INFECTION AND IMMUNITY, (2000 Feb) 68 (2) 732-9.  
Journal code: 0246127. ISSN: 0019-9567.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 200002  
ENTRY DATE: Entered STN: 20000218  
Last Updated on STN: 20000218  
Entered Medline: 20000210

AB Porphyromonas **gingivalis** is a major etiologic agent of periodontitis, a chronic inflammatory disease that ultimately results in the loss of the supporting tissues of the teeth. Previous work has demonstrated the usefulness of avirulent Salmonella enterica serovar Typhimurium strains as antigen delivery systems for protective antigens of pathogens that colonize or cross mucosal surfaces. In this study, we constructed and characterized a recombinant S. enterica serovar Typhimurium avirulent vaccine strain which expresses **hemagglutinin A** and carries no antibiotic resistance markers. **HagA**, a major virulence-associated surface protein, is a potentially useful immunogen that contains an antigenic epitope which, in humans, elicits an immune response that is protective against subsequent colonization by P. **gingivalis**. The **hagA** gene, including its promoter, was cloned into a balanced-lethal Salmonella vector and transferred to the vaccine strain. Heterologous expression of **HagA** was demonstrated in both Escherichia coli JM109 and S. enterica serovar Typhimurium vaccine strain chi4072. The **HagA** epitope was present in its native configuration as determined by immunochemistry and immunoelectron microscopy. Purified recombinant **HagA** was recognized by sera from mice immunized with the S. enterica serovar Typhimurium vaccine strain. The **HagA**-specific antigen of the vaccine was also found to be recognized by serum from a periodontal patient. This vaccine strain, which expresses the functional **hemagglutinin** protein, induces a humoral immune response against **HagA** and may be useful for developing a protective vaccine against periodontal diseases associated with P. **gingivalis**.

L7 ANSWER 5 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 2000:349917 BIOSIS

DOCUMENT NUMBER: PREV200000349917

TITLE: Both **HagA** and HagB **hemagglutinins** of Porphyromonas **gingivalis** act as adhesins for invasion of HCAEC.

AUTHOR(S): Dorn, B. R. (1); Kozarov, E. V. (1); Harris, L. J. (1); Whitlock, J. A. (1); Progulske-Fox, A. (1)

CORPORATE SOURCE: (1) University of Florida, Gainesville, FL USA

SOURCE: Abstracts of the General Meeting of the American Society for Microbiology, (2000) Vol. 100, pp. 70. print.  
Meeting Info.: 100th General Meeting of the American Society for Microbiology Los Angeles, California, USA May 21-25, 2000 American Society for Microbiology  
. ISSN: 1060-2011.

DOCUMENT TYPE: Conference

LANGUAGE: English

SUMMARY LANGUAGE: English

L7 ANSWER 6 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 2000:345558 BIOSIS

DOCUMENT NUMBER: PREV200000345558

TITLE: Receptors for **hemagglutinin A** from Porphyromonas **gingivalis** on human endothelial and epithelial cells.

AUTHOR(S): Kozarov, E. V. (1); Song, Y. H.; Progulske-Fox, A. (1)

CORPORATE SOURCE: (1) University of Florida, Gainesville, FL USA

SOURCE: Abstracts of the General Meeting of the American Society for Microbiology, (2000) Vol. 100, pp. 70. print.  
Meeting Info.: 100th General Meeting of the American

Society for Microbiology Los Angeles, California, USA May  
21-25, 2000 American Society for Microbiology  
. ISSN: 1060-2011.

DOCUMENT TYPE: Conference  
LANGUAGE: English  
SUMMARY LANGUAGE: English

L7 ANSWER 7 OF 38 MEDLINE DUPLICATE 3  
ACCESSION NUMBER: 2000149628 MEDLINE  
DOCUMENT NUMBER: 20149628 PubMed ID: 10685367  
TITLE: Porphyromonas **gingivalis** virulence factors and  
invasion of cells of the cardiovascular system.  
AUTHOR: **Progulske-Fox A**; **Kozarov E**; Dorn B;  
Dunn W Jr; Burks J; Wu Y  
CORPORATE SOURCE: University of Florida, Department of Oral Biology,  
Gainesville 32606, USA.. apfox@dental.ufl.edu  
SOURCE: JOURNAL OF PERIODONTAL RESEARCH, (1999 Oct) 34 (7) 393-9.  
Journal code: 0055107. ISSN: 0022-3484.  
PUB. COUNTRY: Denmark  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Dental Journals; Priority Journals  
ENTRY MONTH: 200003  
ENTRY DATE: Entered STN: 20000330  
Last Updated on STN: 20000330  
Entered Medline: 20000323

AB Our laboratory is interested in the genes and gene products involved in the interactions between Porphyromonas **gingivalis** (Pg) and the host. These interactions may occur in either the periodontal tissues or other non-oral host tissues such as those of the cardiovascular system. We have previously reported the cloning of several genes encoding **hemagglutinins**, surface proteins that interact with the host tissues, and are investigating their roles in the disease process. Primary among these is **HagA**, a very large protein with multiple functional groups that have significant sequence homology to protease genes of this species. Preliminary evidence indicates that an avirulent Salmonella typhimurium strain containing **hagA** is virulent in mice. These data indicate that **HagA** may be a key virulence factor of Pg. Additionally, we are investigating the invasion of primary human coronary artery endothelial cells (HCAEC) by Pg because of the recent epidemiological studies indicating a correlation between periodontal disease (PD) and coronary heart disease (CHD). We found that some, but not all, strains of Pg are able to invade these cells. Scanning electron microscopy of the infected HCAEC demonstrated that the invading organisms initially attached to the host cell surface as aggregates and by a "pedestal"-like structure. By transmission electronmicroscopy it could be seen that internalized bacteria were present within multimembranous compartments localized with rough endoplasmic reticulum. In addition, invasion of the HCAEC by Pg resulted in an increase in the degradation of long-lived cellular proteins. These data indicate that Pg are present within autophagosomes and may use components of the autophagic pathway as a means to survive intracellularly. However, Pg presence within autophagosomes in KB cells could not be observed or detected. It is therefore likely that Pg uses different invasive mechanisms for different host cells. This and the role of **HagA** in invasion is currently being investigated further.

L7 ANSWER 8 OF 38 MEDLINE DUPLICATE 4  
ACCESSION NUMBER: 1998427134 MEDLINE  
DOCUMENT NUMBER: 98427134 PubMed ID: 9746569  
TITLE: The number of direct repeats in **hagA** is variable  
among Porphyromonas **gingivalis** strains.  
AUTHOR: **Kozarov E**; Whitlock J; Dong H; Carrasco E;  
**Progulske-Fox A**

CORPORATE SOURCE: Department of Oral Biology, University of Florida,  
Gainesville, Florida 32610, . USA.kozarov@dental.ufl.edu  
CONTRACT NUMBER: DE07496 (NIDCR)  
SOURCE: INFECTION AND IMMUNITY, (1998 Oct) 66 (10) 4721-5.  
Journal code: 0246127. ISSN: 0019-9567.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199810  
ENTRY DATE: Entered STN: 19990106  
Last Updated on STN: 20000303  
Entered Medline: 19981029

AB The coding sequence for the surface protein **hemagglutinin A** (**HagA**) of *Porphyromonas gingivalis* 381 has previously been shown to contain four direct 1.35-kb repeats, designated repHA. This study was performed to determine if the number of repHA units in **hagA** is consistently 4 or if allelic polymorphism exists among strains and/or upon multiple passage of *P. gingivalis*. To this end, primers which were homologous to the regions directly 5' and 3' of the repeat domain in **hagA** were synthesized. PCR conditions which allowed amplification of the 8.4-kb repeat region between the primers in *P. gingivalis* 381 were established. Genomic DNA templates from 13 other *P. gingivalis* strains and 9 fresh clinical isolates from patients were analyzed under the same conditions as used above. Analysis of these PCR products demonstrated that the strains tested had different numbers (two to four) of repHA units in the respective **hagA** genes. The PCR products of 8.4, 7.0, and 5.7 kb represent four, three, and two repeats, respectively. One strain from each group (381, four repeats; W83, three repeats; and AJW4, two repeats) was also tested to determine if the number of repeats remained invariant upon passaging onto solid medium. No variability in the number of repeats in **hagA** within a strain was detected after 18 passages. *P. gingivalis* 381 was chosen for further testing in a mouse abscess model to determine if conditions of in vivo growth would select for deletions or duplications of the repeated sequences. Five days after infection, no change in the number of repeats was detected in cells recovered from either nonimmunized or preimmunized mice. This data indicates an interstrain variability of the number of repeat units and hence a size variability of the **HagA** protein of *P. gingivalis*, but unlike some surface antigens of other pathogenic species, the number of repeats remains relatively stable given the conditions of growth tested here.

L7 ANSWER 9 OF 38 SCISEARCH COPYRIGHT 2003 ISI (R)  
ACCESSION NUMBER: 1998:411254 SCISEARCH  
THE GENUINE ARTICLE: ZK546  
TITLE: Expression of **hemagglutinin a** from *Porphyromonas gingivalis* in *Escherichia coli* and in a avirulent vaccine strain of *Salmonella typhimurium*  
AUTHOR: Miyashita N (Reprint); Kozarov E V; Burks J N; ProgulskFox A  
CORPORATE SOURCE: UNIV FLORIDA, GAINESVILLE, FL  
COUNTRY OF AUTHOR: USA  
SOURCE: JOURNAL OF DENTAL RESEARCH, (APR 1998) Vol. 77, Sp. iss. B, pp. 1789-1789.  
Publisher: AMER ASSOC DENTAL RESEARCH, 1619 DUKE ST, ALEXANDRIA, VA 22314.  
ISSN: 0022-0345.  
DOCUMENT TYPE: Conference; Journal  
FILE SEGMENT: LIFE; CLIN  
LANGUAGE: English  
REFERENCE COUNT: 0

L7 ANSWER 10 OF 38 SCISEARCH COPYRIGHT 2003 ISI (R)  
 ACCESSION NUMBER: 1998:411253 SCISEARCH  
 THE GENUINE ARTICLE: ZK546  
 TITLE: Porphyromonas **gingivalis** hagB/C  
**hemagglutinin** mutant exhibits reduced invasion of  
 human oral epithelial cells.  
 AUTHOR: Whitlock J A (Reprint); Dorn B R; Burks J N; **Kozarov**  
**E V**; ProgulskeFox A  
 CORPORATE SOURCE: UNIV FLORIDA, GAINESVILLE, FL  
 COUNTRY OF AUTHOR: USA  
 SOURCE: JOURNAL OF DENTAL RESEARCH, (APR 1998) Vol. 77, Sp. iss.  
 B, pp. 1787-1787.  
 Publisher: AMER ASSOC DENTAL RESEARCH, 1619 DUKE ST,  
 ALEXANDRIA, VA 22314.  
 ISSN: 0022-0345.  
 DOCUMENT TYPE: Conference; Journal  
 FILE SEGMENT: LIFE; CLIN  
 LANGUAGE: English  
 REFERENCE COUNT: 0

L7 ANSWER 11 OF 38 MEDLINE DUPLICATE 5  
 ACCESSION NUMBER: 1999138114 MEDLINE  
 DOCUMENT NUMBER: 99138114 PubMed ID: 9972167  
 TITLE: The porphyromonas **gingivalis** prtP/kgp homologue  
 exists as two open reading frames in strain 381.  
 AUTHOR: Han N; Lepine G; Whitlock J; Wojciechowski L;  
**Progulske-Fox A**  
 CORPORATE SOURCE: Department of Oral Biology, University of Florida,  
 Gainesville 32610-0424, USA.  
 CONTRACT NUMBER: DE 00336 (NIDCR)  
 DE 07496 (NIDCR)  
 SOURCE: ORAL DISEASES, (1998 Sep) 4 (3) 170-9.  
 Journal code: 9508565. ISSN: 1354-523X.  
 PUB. COUNTRY: ENGLAND: United Kingdom  
 DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
 LANGUAGE: English  
 FILE SEGMENT: Dental Journals  
 OTHER SOURCE: GENBANK-U68468  
 ENTRY MONTH: 199902  
 ENTRY DATE: Entered STN: 19990301  
 Last Updated on STN: 20000303  
 Entered Medline: 19990218

AB **P. gingivalis** is considered to be a major pathogen of adult  
 periodontitis. Among its cadre of putative virulence factors are  
**hemagglutinins** (adhesins) and proteases. We here report the  
 cloning, sequencing and characterization of two genes, designated kgp(381)  
 and hagD. Kgp(381), an open reading frame (ORF) of 1095 bp encoding a 40.1  
 kda protein, has high homology to the proteolytic domain of cysteine  
 protease/**hemagglutinin** genes. HagD, an ORF of 4077 bp encoding a  
 147.1 kda protein, contains one HArep sequence which establishes it as an  
 additional member of the HArep multigene family. Although similar in  
 sequence to kgp and prtP which were identified from strains HG66 and W12,  
 respectively, the kgp(381)-hagD genes have several characteristics which  
 distinguish them from kgp and prtP. Foremost among these is a single base  
 difference which produces a termination codon and an immediate frame shift  
 resulting in two ORFs in strain 381 as compared to one ORF in strains HG66  
 and W12. In addition, a 172 amino acid sequence near the C-terminal end of  
 hagD has very low identity (20.5-27.8%) to the corresponding region of kgp  
 and prtP. These demonstrate that the homologue of kgp and prtP in strain  
 381 occurs as two separate genes which may genetically separate the  
 adhesive and enzymatic domains of Kgp and PrtP proteins. Reverse  
 polymerase chain reaction (PCR) analysis indicates that hagD expression is  
 regulated by hemin concentration.



L7 ANSWER 12 OF 38 MEDLINE DUPLICATE 6

ACCESSION NUMBER: 97045177 MEDLINE  
DOCUMENT NUMBER: 97045177 PubMed ID: 8890242  
TITLE: The **hemagglutinin** genes **hagB** and **hagC** of *Porphyromonas gingivalis* are transcribed in vivo as shown by use of a new expression vector.  
AUTHOR: Lee S W; Hillman J D; **Progulske-Fox A**  
CORPORATE SOURCE: Department of Oral Biology, College of Dentistry, University of Florida, Gainesville 32610, USA.  
CONTRACT NUMBER: DE00336 (NIDCR)  
DE07496 (NIDCR)  
DE10994 (NIDCR)  
SOURCE: INFECTION AND IMMUNITY, (1996 Nov) 64 (11) 4802-10.  
Journal code: 0246127. ISSN: 0019-9567.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199701  
ENTRY DATE: Entered STN: 19970128  
Last Updated on STN: 20000303  
Entered Medline: 19970106

AB The **hemagglutinin** genes **hagB** and **hagC** of *Porphyromonas gingivalis*, a putative periodontopathic microorganism, have been cloned, sequenced, and characterized. However, the roles of these putative virulence genes have not yet been determined. In this study, an in vivo expression technology vector termed pPGIVET was constructed and used to determine if **hagB** and **hagC** were expressed during an infectious process. We constructed pPGIVET as a conjugative suicide plasmid containing a multiple-cloning site (MCS) upstream of two tandem promoterless reporter genes that encode tetracycline resistance [**tetA(Q)2**] and galactokinase (**galk**). The promoter and a portion of the open reading frame (ORF) of **hagB** were inserted into the MCS in both a positive and a negative orientation relative to the reporter genes. These constructs were conjugated into *P. gingivalis* 381. Southern blot analysis of different transconjugants indicated that Campbell insertions had occurred at the chromosomal **hagB** locus and also at the **hagC** locus, which has high (99%) homology to the ORF of **hagB**. pPGIVET-labeled clones in which the **hag** promoters were positively oriented relative to the reporter genes expressed tetracycline resistance and galactokinase activity in vitro and in vivo at significantly higher levels than did the wild-type strain or clones in which the **hag** promoters were negatively oriented. Expression of tetracycline resistance allowed substantial enrichment of heterodiploids over wild-type cells during a mixed infection in the mouse abscess model. These results indicate that **hagB** and **hagC** are transcriptionally active in vivo and suggested that pPGIVET may be used to isolate *P. gingivalis* genes expressed only during an infectious process.

L7 ANSWER 13 OF 38 MEDLINE DUPLICATE 7

ACCESSION NUMBER: 97047672 MEDLINE  
DOCUMENT NUMBER: 97047672 PubMed ID: 8926061  
TITLE: The **hemagglutinin** gene A (**hagA**) of *Porphyromonas gingivalis* 381 contains four large, contiguous, direct repeats.  
AUTHOR: Han N; Whitlock J; **Progulske-Fox A**  
CORPORATE SOURCE: Department of Oral Biology, University of Florida, Gainesville 32610-0424, USA.. nhan@dental.ufl.edu  
CONTRACT NUMBER: DE 00336 (NIDCR)  
DE 07496 (NIDCR)  
SOURCE: INFECTION AND IMMUNITY, (1996 Oct) 64 (10) 4000-7.  
Journal code: 0246127. ISSN: 0019-9567.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English

FILE SEGMENT: Priority Journals  
OTHER SOURCE: GENBANK-U41807  
ENTRY MONTH: 199611  
ENTRY DATE: Entered STN: 19961219  
Last Updated on STN: 20000303  
Entered Medline: 19961114

AB Porphyromonas **gingivalis** is a gram-negative anaerobic bacterial species strongly associated with adult periodontitis. One of its distinguishing characteristics and putative virulence properties is the ability to agglutinate erythrocytes. We have previously reported the cloning of multiple **hemagglutinin** genes from P. **gingivalis** 381. Subsequent sequencing of clone ST 2 revealed that the cloned fragment contained only an internal portion of the gene which lacked both start and stop codons. We here report the cloning and sequencing of the entire gene, designated **hagA**, as well as its relationship to other genes of this species. By use of inverse PCR technology and the construction of several additional genomic libraries, the complete open reading frame of **hagA** was found to be 7,887 bp in length, encoding a protein of 2,628 amino acids with a molecular mass of 283.3 kDa, which is among the largest genes ever cloned from a prokaryote to date. Within its open reading frame, four large, contiguous, direct repeats (varying from 1,318 to 1,368 bp) were identified. The repeat unit (HAreP), which is assumed to contain the **hemagglutinin** domain, is also present in other recently reported protease and **hemagglutinin** genes in P. **gingivalis**. Thus, we propose that **hagA** and the other genes which share the HAreP sequence form a multigene family with **hagA** as a central member.

L7 ANSWER 14 OF 38 MEDLINE DUPLICATE 8  
ACCESSION NUMBER: 96213011 MEDLINE  
DOCUMENT NUMBER: 96213011 PubMed ID: 8631659  
TITLE: Analysis of the prtP gene encoding porphypain, a cysteine proteinase of Porphyromonas **gingivalis**.  
AUTHOR: Barkocy-Gallagher G A; Han N; Patti J M; Whitlock J; **Progulske-Fox A**; Lantz M S  
CORPORATE SOURCE: Indiana University School of Dentistry, Indianapolis, Indiana 46202, USA.  
CONTRACT NUMBER: DE00336 (NIDCR)  
DE07256-13 (NIDCR)  
DE07496 (NIDCR)  
SOURCE: JOURNAL OF BACTERIOLOGY, (1996 May) 178 (10) 2734-41.  
Journal code: 2985120R. ISSN: 0021-9193.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
OTHER SOURCE: GENBANK-U42210  
ENTRY MONTH: 199607  
ENTRY DATE: Entered STN: 19960715  
Last Updated on STN: 20000303  
Entered Medline: 19960702

AB The cloning and sequencing of the gene encoding porphypain, a cysteine proteinase previously isolated from detergent extracts of the Porphyromonas **gingivalis** W12 cell surface, are described. The prtP gene encoded a unique protein of 1,732 amino acids, including a putative signal sequence for protein secretion. The predicted molecular mass for the mature protein was 186 kDa, which was close to the observed molecular mass of 180 kDa. There was one copy of prtP in the genomes of seven P. **gingivalis** strains examined. The gene was located 5' to a region with a high degree of homology to the insertion element IS1126 in P. **gingivalis** W12. The PrtP protein had regions of high homology to **HagA**, a **hemagglutinin** of P. **gingivalis**, and to several purported proteinases of P. **gingivalis** that have Arg-X specificity. A detailed comparison of genes encoding the latter and

cpgR suggested that rgp-1, prpR1, prtR, agp, cpgR, and possibly prtH were derived from identical genetic loci. Although an rgp-1-like locus was detected in seven *P. gingivalis* strains by Southern blot analyses, agp and cpgR were not detected, not even in the strains from which they were originally isolated. In addition, at least 20 copies of a repeat region common to PrtP, the Rgp-1-like proteins, and **HagA** were observed in each of the seven genomes examined. The repeat region hybridization patterns for strains W83 and W50 were very similar, and they were identical for strains 381 and ATCC 33277, providing further evidence that these strains are closely related genetically.

L7 ANSWER 15 OF 38 MEDLINE DUPLICATE 9

ACCESSION NUMBER: 96178649 MEDLINE

DOCUMENT NUMBER: 96178649 PubMed ID: 8606121

TITLE: Construction and preliminary characterization of three **hemagglutinin** mutants of *Porphyromonas gingivalis*.

AUTHOR: Lepine G; Ellen R P; **Progulske-Fox A**

CORPORATE SOURCE: Department of Oral Biology, College of Dentistry, University of Florida, Gainesville, Florida 32610, USA.

CONTRACT NUMBER: DE00336 (NIDCR)

DE07496 (NIDCR)

SOURCE: INFECTION AND IMMUNITY, (1996 Apr) 64 (4) 1467-72.  
Journal code: 0246127. ISSN: 0019-9567.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199605

ENTRY DATE: Entered STN: 19960531  
Last Updated on STN: 20000303  
Entered Medline: 19960523

AB Targeted insertional mutagenesis was used to construct **hagA**, **hagB**, and **hagC hemagglutinin** mutants of *Porphyromonas gingivalis*. pJRD215-derived plasmids containing tetA(Q)2 and portions of the targeted genes were conjugated into *P. gingivalis*. Interruption of the three loci was confirmed by Southern hybridization, sequencing, reverse transcription-PCR, and microtiter **hemagglutination** assays. No significant differences in hydrophobicity or coadherence to *Actinomyces viscosus* were detected between the mutants and the wild-type strain.

L7 ANSWER 16 OF 38 MEDLINE DUPLICATE 10

ACCESSION NUMBER: 97096936 MEDLINE

DOCUMENT NUMBER: 97096936 PubMed ID: 8941757

TITLE: Duplication and differential expression of **hemagglutinin** genes in *Porphyromonas gingivalis*.

AUTHOR: Lepine G; **Progulske-Fox A**

CORPORATE SOURCE: Department of Oral Biology, College of Dentistry, University of Florida, Gainesville, USA.

CONTRACT NUMBER: DE07496 (NIDCR)

SOURCE: ORAL MICROBIOLOGY AND IMMUNOLOGY, (1996 Apr) 11 (2) 65-78.  
Journal code: 8707451. ISSN: 0902-0055.

PUB. COUNTRY: Denmark

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Dental Journals

OTHER SOURCE: GENBANK-Z27394

ENTRY MONTH: 199701

ENTRY DATE: Entered STN: 19970128  
Last Updated on STN: 20000303  
Entered Medline: 19970114

AB A third **hemagglutinin** gene, defined as **hagC**, was cloned from

Porphyromonas **gingivalis** 381 and sequenced. This gene was found to encode a protein highly homologous (98.6%) to the previously reported HagB **hemagglutinin** protein. The upstream and downstream regions of hagB and hagC were found to share less than 40% homology compared with 99% for their open reading frames. The antigenic relationship between the two **hemagglutinins** was demonstrated by Western blot analysis. When expressed in an in vitro transcription-translation system, both genes encoded a protein with a molecular mass of 49 kDa. As determined by reverse transcription polymerase chain reaction, the steady-state levels of hagB and hagC mRNAs were found to vary according to the growth phase and hemin concentration. The amount of transcripts decreased in hemin-limited conditions or in the absence of hemin. Furthermore, hagB mRNAs were detected in the early logarithmic growth phase compared with the hagC transcripts, which were detected only in the mid-exponential phase of growth.

L7	ANSWER 17 OF 38	MEDLINE	DUPLICATE 11
ACCESSION NUMBER:	96001703	MEDLINE	
DOCUMENT NUMBER:	96001703	PubMed ID: 7502765	
TITLE:	Expression and immunogenicity of a cloned Porphyromonas <b>gingivalis</b> <b>hemagglutinin</b> in Salmonella typhimurium.		
AUTHOR:	Dusek D M; <b>Progulske-Fox A</b> ; Brown T A		
CORPORATE SOURCE:	Department of Oral Biology, University of Florida, Gainesville, USA.		
CONTRACT NUMBER:	DE00236 (NIDCR)		
	DE07117 (NIDCR)		
	DE07496 (NIDCR)		
	+		
SOURCE:	ADVANCES IN EXPERIMENTAL MEDICINE AND BIOLOGY, (1995) 371B 1119-21.		
	Journal code: 0121103. ISSN: 0065-2598.		
PUB. COUNTRY:	United States		
DOCUMENT TYPE:	Journal; Article; (JOURNAL ARTICLE)		
LANGUAGE:	English		
FILE SEGMENT:	Priority Journals		
ENTRY MONTH:	199601		
ENTRY DATE:	Entered STN: 19960217		
	Last Updated on STN: 20000303		
	Entered Medline: 19960116		

  

L7	ANSWER 18 OF 38	MEDLINE	DUPLICATE 12
ACCESSION NUMBER:	96179248	MEDLINE	
DOCUMENT NUMBER:	96179248	PubMed ID: 8596675	
TITLE:	The cloning, expression and sequence analysis of a second Porphyromonas <b>gingivalis</b> gene that codes for a protein involved in <b>hemagglutination</b> .		
AUTHOR:	<b>Progulske-Fox A</b> ; Tumwasorn S; Lepine G; Whitlock J; Savett D; Ferretti J J; Banas J A		
CORPORATE SOURCE:	Department of Oral biology, College of Dentistry, University of Florida, Gainesville, USA.		
CONTRACT NUMBER:	DE00336 (NIDCR)		
	DE005545 (NIDCR)		
	DE007496 (NIDCR)		
SOURCE:	ORAL MICROBIOLOGY AND IMMUNOLOGY, (1995 Oct) 10 (5) 311-8.		
	Journal code: 8707451. ISSN: 0902-0055.		
PUB. COUNTRY:	Denmark		
DOCUMENT TYPE:	Journal; Article; (JOURNAL ARTICLE)		
LANGUAGE:	English		
FILE SEGMENT:	Dental Journals		
OTHER SOURCE:	GENBANK-Z35494		
ENTRY MONTH:	199604		
ENTRY DATE:	Entered STN: 19960424		
	Last Updated on STN: 20000303		

Entered Medline: 19960418

AB It has been suggested that *Porphyromonas gingivalis* may possess more than one **hemagglutinin**. We have previously reported the cloning of a gene (**hagA**) that encodes a **hemagglutinin**. In this study we report the cloning, characterization, and sequencing of a second gene (**hagB**) that encodes a protein that also appears to be involved in **hemagglutination**. Antiserum to the clone (ST 7) was found to inhibit **hemagglutination** by *P. gingivalis* 381, and **hemagglutinating** inhibition activity of anti-*P. gingivalis* antiserum was reduced by adsorption of the antiserum with cells of clone ST 7. Restriction mapping and Southern analysis indicates there is little or no DNA homology between this cloned 4.8-kb HindIII DNA fragment and a cloned **hemagglutinin** gene we have previously described. Minicell analysis of the cloned *P. gingivalis* chromosomal DNA fragment revealed that the major gene product is a 49-kDa protein. Immunoaffinity chromatography using purified rabbit immunoglobulin G against the cloned protein resulted in the purification of a major reactive 49- to 50-kDa protein from a *P. gingivalis* cell lysate. Nucleotide sequence analysis revealed the **hagB** open reading frame to be 1053 nucleotides in length with a mol% G+C of 59.9% coding for a protein of 350 residues with a calculated molecular weight of 39.375 kDa. This protein was also determined to be basic and hydrophilic and to contain a potential signal peptide. Comparison of both the nucleotide and derived amino acid sequences with computer-based databases did not reveal any significant homologies between **hagB** and any other previously sequenced genes.

L7 ANSWER 19 OF 38 MEDLINE DUPLICATE 13  
ACCESSION NUMBER: 95372118 MEDLINE  
DOCUMENT NUMBER: 95372118 PubMed ID: 7644268  
TITLE: Restriction fragment length polymorphism analysis of two **hemagglutinin** loci, serotyping and agglutinating activity of *Porphyromonas gingivalis* isolates.  
AUTHOR: Savett D A; **Progulske-Fox A**  
CORPORATE SOURCE: Department of Oral Biology, College of Dentistry, University of Florida, Gainesville 32610-0424, USA.  
CONTRACT NUMBER: DE07496 (NIDCR)  
SOURCE: ORAL MICROBIOLOGY AND IMMUNOLOGY, (1995 Feb) 10 (1) 1-7. Journal code: 8707451. ISSN: 0902-0055.  
PUB. COUNTRY: Denmark  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Dental Journals  
ENTRY MONTH: 199509  
ENTRY DATE: Entered STN: 19950930  
Last Updated on STN: 20000303  
Entered Medline: 19950918

AB Restriction fragment length polymorphisms (RFLPs) of two **hemagglutinin** loci were analyzed in 36 *Porphyromonas gingivalis* isolates from human and monkey origins using portions of **hagA** and **hagB** as probes. The *P. gingivalis* strains were differentiated into 9 RFLP groups based on the heterogeneity of the **hagA** locus and 10 different groups based on hybridization with **hagB**. Homology to **hagA** was detected in all human derived and all but three monkey derived strains. All *P. gingivalis* isolates exhibited DNA homologous to **hagB**. Multiple alleles of the **hemagglutinin** genes were detected for most *P. gingivalis* strains. No DNA homologous to either **hemagglutinin** gene could be detected in 6 other bacterial species tested. Serotyping and **hemagglutination** titers of each *P. gingivalis* isolate were obtained in an attempt to establish a correlation between these pheno-typic parameters and RFLP group. Although no correlations were found with these parameters, a correlation between RFLP group and invasiveness in the mouse abscess model was noted.

L7 ANSWER 20 OF 38

MEDLINE

DUPLICATE 14

ACCESSION NUMBER: 94222526 MEDLINE  
DOCUMENT NUMBER: 94222526 PubMed ID: 8168925  
TITLE: Systemic and mucosal immune responses in mice orally  
immunized with avirulent *Salmonella typhimurium* expressing  
a cloned *Porphyromonas gingivalis*  
*hemagglutinin*.  
AUTHOR: Dusek D M; **Progulske-Fox A**; Brown T A  
CORPORATE SOURCE: Department of Oral Biology, University of Florida,  
Gainesville 32610.  
CONTRACT NUMBER: DE 07117 (NIDCR)  
DE 07496 (NIDCR)  
RCDA DE 00236 (NIDCR)  
SOURCE: INFECTION AND IMMUNITY, (1994 May) 62 (5) 1652-7.  
Journal code: 0246127. ISSN: 0019-9567.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199406  
ENTRY DATE: Entered STN: 19940613  
Last Updated on STN: 20000303  
Entered Medline: 19940602

AB *Porphyromonas gingivalis* produces a variety of virulence factors that may have a function in the periodontal disease process. Determination of the role of these various factors in pathogenesis and identification of a means for protecting the host from the destructive effects of this organism are areas of vigorous investigation. In this study we demonstrate the potential of avirulent *Salmonella typhimurium* strains to stimulate a specific systemic and mucosal immune response to a cloned *P. gingivalis* *hemagglutinin* (HagB). An avirulent strain of *S. typhimurium*, chi 4072, expressing the hagB gene of *P. gingivalis* 381 on the plasmid pDMD1 was intragastrically administered to BALB/c mice. These mice mounted a serum immunoglobulin G (IgG) and IgA primary response against the hagB gene product and a mucosal immune response as measured by evaluation of saliva. IgA antibodies were also detected in bile. These results demonstrate the feasibility of using attenuated *S. typhimurium* strains as carriers of *P. gingivalis* virulence factors for subsequent evaluation of the systemic and mucosal immune response against these antigens. This system will provide a means for evaluating the virulence factors of *P. gingivalis* for their suitability in the construction of potential vaccines.

L7 ANSWER 21 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 1994:330724 BIOSIS  
DOCUMENT NUMBER: PREV199497343724  
TITLE: Interactions of *Porphyromonas gingivalis* with  
oral epithelial cells.  
AUTHOR(S): Emory, S. (1); Duncan, M.; Lepine, G.; Han, N.;  
**Progulske-Fox, A.**  
CORPORATE SOURCE: (1) Forsyth Dental Center, Boston, MA USA  
SOURCE: Abstracts of the General Meeting of the American Society  
for Microbiology, (1994) Vol. 94, No. 0, pp. 116.  
Meeting Info.: 94th General Meeting of the American Society  
for Microbiology Las Vegas, Nevada, USA May 23-27, 1994  
ISSN: 1060-2011.  
DOCUMENT TYPE: Conference  
LANGUAGE: English

L7 ANSWER 22 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 1994:87296 BIOSIS  
DOCUMENT NUMBER: PREV199497100296  
TITLE: Molecular biology.  
AUTHOR(S): Lepine, Guylaine (1); **Progulske-Fox, Ann**

CORPORATE SOURCE: (1) Dep. Oral Biology, Univ. Fla., Gainesville, FL USA  
SOURCE: Shah, H. N. [Editor]. (1993) pp. 293-319. Biology of the  
species Porphyromonas gingivalis.  
Publisher: CRC Press, Inc. Boca Raton, Florida, USA.  
ISBN: 0-8493-6648-8.  
DOCUMENT TYPE: Book  
LANGUAGE: English

L7 ANSWER 23 OF 38 MEDLINE DUPLICATE 15  
ACCESSION NUMBER: 93162835 MEDLINE  
DOCUMENT NUMBER: 93162835 PubMed ID: 8381773  
TITLE: Isolation and characterization of a cloned Porphyromonas  
**gingivalis hemagglutinin** from an  
avirulent strain of Salmonella typhimurium.  
AUTHOR: Dusek D M; **Progulske-Fox A**; Whitlock J; Brown T A  
CORPORATE SOURCE: Department of Oral Biology, University of Florida,  
Gainesville 32610.  
CONTRACT NUMBER: DE 00236 (NIDCR)  
DE 07496 (NIDCR)  
DE07117 (NIDCR)  
SOURCE: INFECTION AND IMMUNITY, (1993 Mar) 61 (3) 940-6.  
Journal code: 0246127. ISSN: 0019-9567.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199303  
ENTRY DATE: Entered STN: 19930402  
Last Updated on STN: 20000303  
Entered Medline: 19930318

AB Identification of surface macromolecules of Porphyromonas  
**gingivalis** that act as virulence factors in periodontal disease  
has important implications for studying host-parasite interactions as well  
as for potential vaccine development. The objective of this study was to  
determine whether a cloned, P. **gingivalis hemagglutinin**  
gene could be expressed in an intact form in an avirulent Salmonella  
typhimurium vaccine construct and to characterize the recombinant protein.  
The recombinant protein was purified from the vaccine strain,  
characterized, and tested for biological activity as a competitive  
inhibitor of **hemagglutination**. Cells of S. typhimurium  
SL3261/pST7 grown in Luria broth were broken by sonic disruption and  
fractionated. The purified recombinant protein was found to inhibit  
**hemagglutination** of erythrocytes by whole P. **gingivalis**  
cells. The same purified protein was analyzed for its N-terminal amino  
acid sequence and amino acid composition and found to match that predicted  
from the nucleotide sequence of the cloned gene. These results indicate  
that a surface macromolecule of P. **gingivalis** can be expressed  
in an intact and biologically active form in a Salmonella carrier strain.

L7 ANSWER 24 OF 38 MEDLINE DUPLICATE 16  
ACCESSION NUMBER: 94087471 MEDLINE  
DOCUMENT NUMBER: 94087471 PubMed ID: 8263716  
TITLE: Molecular characterization of **hemagglutinin** genes  
of periodontopathic bacteria.  
AUTHOR: **Progulske-Fox A**; Rao V; Han N; Lepine G; Witlock  
J; Lantz M  
CORPORATE SOURCE: Department of Oral Biology, University of Florida,  
Gainesville 32610-0424.  
SOURCE: JOURNAL OF PERIODONTAL RESEARCH, (1993 Nov) 28 (6 Pt 2)  
473-4.  
Journal code: 0055107. ISSN: 0022-3484.  
PUB. COUNTRY: Denmark  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English

FILE SEGMENT: Dental Journals; Priority Journals  
ENTRY MONTH: 199401  
ENTRY DATE: Entered STN: 19940209  
Last Updated on STN: 19990129  
Entered Medline: 19940121

L7 ANSWER 25 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1993:201532 BIOSIS  
DOCUMENT NUMBER: PREV199344097782  
TITLE: DNA sequence analysis of a gene encoding a putative  
fibrinogen and fibronectin binding protein from  
Porphyromonas **gingivalis**.  
AUTHOR(S): Han, N. (1); Whitlock, J.; Patti, J. M.; Lantz, M. S.;  
**Progulske-Fox, A.**  
CORPORATE SOURCE: (1) Univ. Fla., Gainesville, FL USA  
SOURCE: Journal of Dental Research, (1993) Vol. 72, No. ABSTR.  
SPEC. ISSUE, pp. 156.  
Meeting Info.: Joint Meeting of the International  
Association for Dental Research, the American Association  
of Dental Research and the Canadian Association of Dental  
Research Chicago, Illinois, USA March 10-14, 1993  
ISSN: 0022-0345.  
DOCUMENT TYPE: Conference  
LANGUAGE: English

L7 ANSWER 26 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1993:201335 BIOSIS  
DOCUMENT NUMBER: PREV199344097585  
TITLE: Isogenic mutations in two **hemagglutinin** genes of  
Porphyromonas **gingivalis**.  
AUTHOR(S): Lepine, G.; **Progulske-Fox, A.**  
CORPORATE SOURCE: Dep. Oral Biol., Univ. Fla., Gainesville, FL USA  
SOURCE: Journal of Dental Research, (1993) Vol. 72, No. ABSTR.  
SPEC. ISSUE, pp. 118.  
Meeting Info.: Joint Meeting of the International  
Association for Dental Research, the American Association  
of Dental Research and the Canadian Association of Dental  
Research Chicago, Illinois, USA March 10-14, 1993  
ISSN: 0022-0345.  
DOCUMENT TYPE: Conference  
LANGUAGE: English

L7 ANSWER 27 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1993:357458 BIOSIS  
DOCUMENT NUMBER: PREV199345040883  
TITLE: Cloning and characterization of a third Porphyromonas  
**gingivalis hemagglutinin** gene.  
AUTHOR(S): Lepine, G.; **Progulske-Fox, A.**  
CORPORATE SOURCE: Univ. Fla., Gainesville, FL USA  
SOURCE: Abstracts of the General Meeting of the American Society  
for Microbiology, (1993) Vol. 93, No. 0, pp. 116.  
Meeting Info.: 93rd General Meeting of the American Society  
for Microbiology Atlanta, Georgia, USA May 16-20, 1993  
ISSN: 1060-2011.  
DOCUMENT TYPE: Conference  
LANGUAGE: English

L7 ANSWER 28 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1993:357454 BIOSIS  
DOCUMENT NUMBER: PREV199345040879  
TITLE: Serotype and **hemagglutination** titers of  
Porphyromonas **gingivalis**.  
AUTHOR(S): Savett, D. A.; Lepine, G.; **Progulske-Fox, A.**  
CORPORATE SOURCE: Univ. Fla., Gainesville, FL USA



SOURCE: Abstracts of the General Meeting of the American Society  
for Microbiology, (1993) Vol. 93, No. 0, pp. 115.  
Meeting Info.: 93rd General Meeting of the American Society  
for Microbiology Atlanta, Georgia, USA May 16-20, 1993  
ISSN: 1060-2011.  
DOCUMENT TYPE: Conference  
LANGUAGE: English

L7 ANSWER 29 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1992:401937 BIOSIS  
DOCUMENT NUMBER: BR43:57812  
TITLE: IMMUNE RESPONSE OF MICE ORALLY IMMUNIZED WITH CLONED  
PORPHYROMONAS-**GINGIVALIS** **HEMAGGLUTININ**.  
AUTHOR(S): DUSEK D M; **PROGULSKE-FOX A**; BROWN T A  
CORPORATE SOURCE: DEP. ORAL BIOL., UNIV. FLA., GAINESVILLE, FLA.  
SOURCE: JOINT MEETING OF THE 70TH GENERAL MEETING OF THE  
INTERNATIONAL ASSOCIATION FOR DENTAL RESEARCH (IADR), 40TH  
ANNUAL MEETING OF THE BRITISH DIVISION OF THE IADR, 1992  
ANNUAL MEETING OF THE CONTINENTAL EUROPEAN DIVISION OF THE  
IADR, 8TH ANNUAL MEETING OF THE IRISH DIVISION OF THE IADR,  
AND THE 75TH ANNUAL MEETING OF THE SCANDINAVIAN ASSOCIATION  
FOR DENTAL RESEARCH, GLASGOW, SCOTLAND, UK, JULY 1-4, 1992.  
J DENT RES, (1992) 71 (SPEC ISSUE), 756.  
CODEN: JDREAF. ISSN: 0022-0345.  
DOCUMENT TYPE: Conference  
FILE SEGMENT: BR; OLD  
LANGUAGE: English

L7 ANSWER 30 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1992:382047 BIOSIS  
DOCUMENT NUMBER: BR43:48997  
TITLE: NUCLEIC ACID SEQUENCE ANALYSIS OF GENES ENCODING  
**HEMAGGLUTININ** AND FIBRINOGEN BINDING ACTIVITIES OF  
PORPHYROMONAS-**GINGIVALIS**.  
AUTHOR(S): HAN N; WHITLOCK J; PATTI J M; LANTZ M S; **PROGULSKE-FOX**  
**A**  
CORPORATE SOURCE: UNIV. FLA., GAINESVILLE, FLA.  
SOURCE: JOINT MEETING OF THE 70TH GENERAL MEETING OF THE  
INTERNATIONAL ASSOCIATION FOR DENTAL RESEARCH (IADR), 40TH  
ANNUAL MEETING OF THE BRITISH DIVISION OF THE IADR, 1992  
ANNUAL MEETING OF THE CONTINENTAL EUROPEAN DIVISION OF THE  
IADR, 8TH ANNUAL MEETING OF THE IRISH DIVISION OF THE IADR  
AND THE 75TH ANNUAL MEETING OF THE SCANDINAVIAN ASSOCIATION  
FOR DENTAL RESEARCH, GLASGOW, SCOTLAND, UK, JULY 1-4, 1992.  
J DENT RES, (1992) 71 (SPEC ISSUE), 530.  
CODEN: JDREAF. ISSN: 0022-0345.  
DOCUMENT TYPE: Conference  
FILE SEGMENT: BR; OLD  
LANGUAGE: English

L7 ANSWER 31 OF 38 MEDLINE DUPLICATE 17  
ACCESSION NUMBER: 92219240 MEDLINE  
DOCUMENT NUMBER: 92219240 PubMed ID: 1313881  
TITLE: Evidence for independent molecular identity and functional  
interaction of the haemagglutinin and cysteine proteinase  
(gingivain) of *Porphyromonas gingivalis*.  
AUTHOR: Shah H N; Gharbia S E; **Progulske-Fox A**;  
Brocklehurst K  
CORPORATE SOURCE: Department of Oral Microbiology, London Hospital Medical  
College, University of London, Whitechapel.  
SOURCE: JOURNAL OF MEDICAL MICROBIOLOGY, (1992 Apr) 36 (4) 239-44.  
Journal code: 0224131. ISSN: 0022-2615.  
PUB. COUNTRY: ENGLAND: United Kingdom  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199205  
ENTRY DATE: Entered STN: 19920529  
Last Updated on STN: 20000303  
Entered Medline: 19920514

AB The sequence of events involved in haemagglutination and lysis of erythrocytes by washed cells, vesicles and the culture supernate of *Porphyromonas gingivalis* strain W83 was monitored by <sup>51</sup>Cr release and transmission electronmicroscopy. All preparations, except capsular material and lipopolysaccharide, caused haemagglutination and, by a slow process of attachment and specific attack on the surface structures of the red blood cells, produced minute pores and eventual leakage of cellular contents. N-acetylglucosamine, N-acetylgalactosamine and several other sugars such as glucose and sucrose had no effect on haemagglutination. Antiserum raised against a cloned haemagglutinin of *P. gingivalis* strain 381 inhibited the activity of strain W83 cells, vesicles and supernate. The antiserum-neutralised supernate lost 70-80% of its hydrolytic activity towards alpha-N-benzoyl-L-arginine-4-nitroanilide but the residual activity behaved in a manner similar to the native supernate in that it was completely inhibited by the addition of 2,2'-dipyridyl disulphide and was fully restored upon addition of a low-Mr mercaptan. Binding of the antiserum to the haemagglutinin epitope of *P. gingivalis* still permitted titration of the active centre cysteinyl thiol group of the proteinase. Purified gingivain caused lysis of erythrocytes and was not neutralised by antiserum to the haemagglutinin. These results suggest that, although the haemagglutinin and gingivain are probably separate molecules, they are closely associated on the outer membrane of *P. gingivalis* and may be functionally related.

L7 ANSWER 32 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 1991:241745 BIOSIS

DOCUMENT NUMBER: BR40:115910

TITLE: OCCURRENCE OF PORPHYROMONAS-**GINGIVALIS** 381  
**HEMAGGLUTININ** GENES IN OTHER PORPHYROMONAS-  
**GINGIVALIS** STRAINS AND ORAL BACTERIAL SPECIES.

AUTHOR(S): LEPINE G; **PROGULSKE-FOX A**

CORPORATE SOURCE: DEP. ORAL BIOL., UNIV. FLA., GAINESVILLE, FLA.

SOURCE: 69TH GENERAL SESSION OF THE INTERNATIONAL ASSOCIATION FOR DENTAL RESEARCH, 20TH ANNUAL SESSION OF THE AMERICAN ASSOCIATION FOR DENTAL RESEARCH, AND THE 12TH ANNUAL SESSION OF THE MEXICAN DIVISION OF THE IADR (INTERNATIONAL ASSOCIATION FOR DENTAL RESEARCH), ACAPULCO, MEXICO, APRIL 17-21, 1991. J DENT RES, (1991) 70 (SPEC ISSUE APRIL), 582. CODEN: JDREAF. ISSN: 0022-0345.

DOCUMENT TYPE: Conference

FILE SEGMENT: BR; OLD

LANGUAGE: English

L7 ANSWER 33 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 1991:240846 BIOSIS

DOCUMENT NUMBER: BR40:115011

TITLE: PURIFICATION AND CHARACTERIZATION OF A PUTATIVE  
**HEMAGGLUTININ** OF PORPHYROMONAS-**GINGIVALIS**

AUTHOR(S): DUSEK D M; WHITLOCK J; **PROGULSKE-FOX A**; BROWN T A

CORPORATE SOURCE: UNIV. FLA., USA.

SOURCE: 69TH GENERAL SESSION OF THE INTERNATIONAL ASSOCIATION FOR DENTAL RESEARCH, 20TH ANNUAL SESSION OF THE AMERICAN ASSOCIATION FOR DENTAL RESEARCH, AND THE 12TH ANNUAL SESSION OF THE MEXICAN DIVISION OF THE IADR (INTERNATIONAL ASSOCIATION FOR DENTAL RESEARCH), ACAPULCO, MEXICO, APRIL 17-21, 1991. J DENT RES, (1991) 70 (SPEC ISSUE APRIL), 437.

CODEN: JDREAF. ISSN: 0022-0345.

DOCUMENT TYPE: Conference  
FILE SEGMENT: BR; OLD  
LANGUAGE: English

L7 ANSWER 34 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1990:365437 BIOSIS  
DOCUMENT NUMBER: BR39:49913  
TITLE: THE CLONING AND CHARACTERIZATION OF A SECOND PORPHYROMONAS-  
**GINGIVALIS HEMAGGLUTININ** GENE.  
AUTHOR(S): TUMWASORN S; LEPINE G; AVAMPATO J; BANAS J; SAVETT D;  
**PROGULSKE-FOX A**  
CORPORATE SOURCE: UNIV. FLA., GAINESVILLE, FLA.  
SOURCE: 90TH ANNUAL MEETING OF THE AMERICAN SOCIETY FOR  
MICROBIOLOGY 1990, ANAHEIM, CALIFORNIA, USA, MAY 13-17,  
1990. ABSTR ANNU MEET AM SOC MICROBIOL, (1990) 90 (0), 93.  
CODEN: ASMACK. ISSN: 0094-8519.  
DOCUMENT TYPE: Conference  
FILE SEGMENT: BR; OLD  
LANGUAGE: English

L7 ANSWER 35 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1989:259325 BIOSIS  
DOCUMENT NUMBER: BR36:126549  
TITLE: IDENTIFICATION OF ANTIGENS OF BACTEROIDES-  
**GINGIVALIS** WHICH HAVE **HEMAGGLUTINATION**  
ACTIVITY.  
AUTHOR(S): SAVETT D; AVAMPATO J; MOORE L; **PROGULSKE-FOX A**  
CORPORATE SOURCE: UNIV. FLA., GAINESVILLE, FLA.  
SOURCE: 18TH ANNUAL SESSION OF THE AMERICAN ASSOCIATION FOR DENTAL  
RESEARCH, SAN FRANCISCO, CALIFORNIA, USA, MARCH 15-19,  
1989. J DENT RES, (1989) 68 (SPEC ISSUE), 356.  
CODEN: JDREAF. ISSN: 0022-0345.  
DOCUMENT TYPE: Conference  
FILE SEGMENT: BR; OLD  
LANGUAGE: English

L7 ANSWER 36 OF 38 MEDLINE DUPLICATE 18  
ACCESSION NUMBER: 90310556 MEDLINE  
DOCUMENT NUMBER: 90310556 PubMed ID: 2700777  
TITLE: The expression and function of a Bacteroides  
**gingivalis hemagglutinin** gene in  
Escherichia coli.  
AUTHOR: **Progulske-Fox A**; Tumwasorn S; Holt S C  
SOURCE: ORAL MICROBIOLOGY AND IMMUNOLOGY, (1989 Sep) 4 (3) 121-31.  
Journal code: 8707451. ISSN: 0902-0055.  
PUB. COUNTRY: Denmark  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Dental Journals  
ENTRY MONTH: 199008  
ENTRY DATE: Entered STN: 19900921  
Last Updated on STN: 19900921  
Entered Medline: 19900813

AB Eight Escherichia coli JM 109 transformants generated from a clone bank of  
Bacteroides **gingivalis** 381 genomic DNA, were found to express B.  
**gingivalis** antigens. Quantitation of antigen expression by ELISA  
indicated that isopropyl-beta-D-thiogalactopyranoside (IPTG) was not  
necessary for antigen expression for any of the clones but that expression  
in 2 of the clones, ST 2 and ST 3, was increased in cells grown in the  
presence of IPTG. Western blot analysis revealed that the expressed  
protein of clone ST 2 has a molecular weight of 125,000 Dal. and that  
clone ST 3 contains multiple bands of 30 to 50 kdal which react with the  
anti-B. **gingivalis** antiserum. Three of the transformants were

found to agglutinate sheep erythrocytes. Polyclonal monospecific antiserum to one of the transformants, clone ST 2, was found to react to 2 major bands of MWs 43,000 and 38,000 and minor bands of 115,000, 105,000, 32,000, and 30,000 Dal. present in *B. gingivalis* cell lysate preparations. Adsorption of anti *B. gingivalis* antiserum with cells of clone ST 2 reduced the hemagglutination inhibition activity of the antiserum 4-fold whereas antiserum to the clone itself inhibited *B. gingivalis* hemagglutination at a titer of 8 times that of normal rabbit serum. Immunoelectronmicroscopic studies using the antiserum to clone ST 2 indicate that the product of the cloned gene (hemagglutinin) is located on the *B. gingivalis* cell surface. A restriction map generated of the cloned *B. gingivalis* DNA fragment confirms the insert to be 3.2 kbases and indicates the possibility of a repeated sequence in the fragment.

L7 ANSWER 37 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1988:260098 BIOSIS  
DOCUMENT NUMBER: BR34:131128  
TITLE: EXPRESSION OF BACTEROIDES-**GINGIVALIS**  
**HEMAGGLUTININS** IN ESCHERICHIA-COLI.  
AUTHOR(S): TUMWASORN S; **PROGULSKE A**  
CORPORATE SOURCE: DEP. ORAL BIOL., PERIODONTAL DIS. RES. CENT., UNIV.  
FLORIDA, GAINESVILLE, FLA.  
SOURCE: 66TH GENERAL SESSION OF THE INTERNATIONAL ASSOCIATION FOR  
DENTAL RESEARCH, 17TH ANNUAL SESSION OF THE AMERICAN  
ASSOCIATION FOR DENTAL RESEARCH, AND 12TH ANNUAL MEETING OF  
THE CANADIAN ASSOCIATION FOR DENTAL RESEARCH, MONTREAL,  
QUEBEC, CANADA, MARCH 9-13, 1988. J DENT RES, (1988) 67  
(SPEC ISSUE MAR ), 368.  
CODEN: JDREAF. ISSN: 0022-0345.  
DOCUMENT TYPE: Conference  
FILE SEGMENT: BR; OLD  
LANGUAGE: English

L7 ANSWER 38 OF 38 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 1988:342577 BIOSIS  
DOCUMENT NUMBER: BR35:37419  
TITLE: THE EXPRESSION OF A BACTEROIDES-**GINGIVALIS**  
**HEMAGGLUTININ** GENE IN ESCHERICHIA-COLI.  
AUTHOR(S): TUMWASORN S; AVAMPATO J; SAVETT D; HOLT S C; **PROGULSKE A**  
CORPORATE SOURCE: UNIV. TEXAS, SAN ANTONIO, TEX.  
SOURCE: ANNUAL MEETING OF THE AMERICAN SOCIETY FOR MICROBIOLOGY,  
MIAMI BEACH, FLORIDA, USA, MAY 8-13, 1988. ABSTR ANNU MEET  
AM SOC MICROBIOL, (1988) 88 (0), 94.  
CODEN: ASMACK. ISSN: 0094-8519.  
DOCUMENT TYPE: Conference  
FILE SEGMENT: BR; OLD  
LANGUAGE: English

# WEST Search History

DATE: Tuesday, March 04, 2003

<u>Set Name</u>	<u>Query</u>	<u>Hit Count</u>	<u>Set Name</u>
side by side			result set
<i>DB=USPT; PLUR=YES; OP=OR</i>			
L47	144 and (protease or proteoly\$)	35	L47
L46	L43 and (bacterium or bacteria).ab.	0	L46
L45	143 and bacteri\$.ti.	0	L45
L44	L43 and (bacterium or bacteria)	52	L44
L43	129 and (\$carcinoma or melanoma or sarcoma or cancer\$ or neoplas\$ or tumor\$ or metast\$).clm.	118	L43
<i>DB=USPT,PGPB,JPAB,EPAB,DWPI; PLUR=YES; OP=OR</i>			
L42	L41 and (bacteri\$)	154	L42
L41	129 and (\$carcinoma or melanoma or sarcoma or cancer\$ or neoplas\$ or tumor\$ or metast\$).clm.	226	L41
L40	122 and (\$carcinoma or melanoma or sarcoma or cancer\$ or neoplas\$ or tumor\$ or metast\$).clm.	17	L40
L39	L38 and (\$carcinoma or melanoma or sarcoma or cancer\$ or neoplas\$ or tumor\$ or metast\$)	0	L39
L38	5830710.pn.	2	L38
L37	(Prtp or dentilisin).ab.	8	L37
L36	L35 and (124 or 18)	2	L36
L35	gingipains.ab.	19	L35
L34	L33 and (18 or 124)	40	L34
L33	((424/197.11)!.CCLS. )	286	L33
L32	L30 and (bacter\$).ab.	12	L32
L31	L30 and 19	1	L31
L30	L29 and (124 or 18)	300	L30
L29	((424/94.1  424/94.2  424/94.21  424/94.3  424/94.4  424/94.5  424/94.6  424/94.61  424/94.62  424/94.63  424/94.64  424/94.65  424/94.66  424/94.67)!.CCLS. )	2936	L29
L28	L27 and (124 or 18)	0	L28
L27	L26 and bacter\$.ab.	51	L27
L26	((424/94.1)!.CCLS. )	683	L26
L25	L24 and 122	1	L25
L24	(angiogen\$ or antiangiogen\$ or neovas\$)	14274	L24
L23	L22 and 18	15	L23
L22	((424/190.1)!.CCLS. )	401	L22
L21	L20 and (bacteri\$).ab.	645	L21
L20	enzyme.ab. and 18	4966	L20

L19	L18 and l8	0	L19
L18	(protease adj bacterium).ab.	11	L18
L17	L16 and l8	0	L17
L16	("bacterial protease").ab.	82	L16
L15	L13 and (angiogen\$ or antiangiogen\$ or neovas\$)	5	L15
L14	L13 and gingivalis	0	L14
L13	L12 and @py<=2001	101	L13
L12	L11 and (bacterium or bacterial).ab.	126	L12
L11	l8 and (protease or aminopept\$).ab.	1400	L11
L10	l8 and l9	21	L10
L9	gingivalis	688	L9
L8	(\$carcinoma or melanoma or sarcoma or cancer\$ or neoplas\$ or tumor\$ or metast\$).ab.	88527	L8
L7	l6 and (angiogen\$ or neovascu\$ or angio\$ or \$carcinoma or cancer\$ or neoplas\$ or tumor\$ or metast\$)	2	L7
L6	(Kozarov-E\$ or Progulske-A\$ or Progulske-fox-a\$).in.	8	L6
L5	l1 and (angiogen\$ or neovascu\$ or antiangio\$)	21	L5
L4	l2 and (\$carcinoma or cancer\$ or neoplas\$ or tumor\$ or metast\$).ab.	20	L4
L3	L2 and gingivalis	0	L3
L2	L1 and (angio\$ or neovas\$ or antiangio\$).ab.	42	L2
L1	(bacter\$).ab. and (protease or proteoly\$).ab.	1977	L1

END OF SEARCH HISTORY